of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this document. Any objection received in response to the regulation may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 177

Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Director, Center for Food Safety and Applied Nutrition, 21 CFR part 177 is amended as follows:

PART 177—INDIRECT FOOD ADDITIVES: POLYMERS

1. The authority citation for 21 CFR part 177 continues to read as follows:

Authority: 21 U.S.C. 321, 342, 348, 379e.

2. Section 177.1520 is amended in the table in paragraph (c) by revising item ''6.'' under the heading ''Density'' to read as follows:

§ 177.1520 Olefin polymers.

* * * * *

(c) * * *

Olefin polymers	Density	Melting Point (MP) or softening point (SP) (*Degrees Centigrade*)	Maximum extractable fraction (expressed as percent by weight of the polymer) in *N*-hexane at specified temperatures	Maximum soluble fraction (expressed as percent by weight of polymer) in xylene at specified temperatures
	* *	* * *		
6. Ethylene-maleic anhydride copolymers described in paragraph (a)(6) of this section for use as the adhesive component in multilaminate structures, or as the sealant layer in flexible packaging, in contact with food at temperatures not exceeding 49 °C (120 °F)	0.92 or greater		1.36 pct at 50 °C.	2.28 pct at 25 °C

* * * * *

Dated: May 5, 1999.

L. Robert Lake,

Director, Office of Policy, Planning and Strategic Initiatives, Center for Food Safety and Applied Nutrition.

[FR Doc. 99–12962 Filed 5–21–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 522

Implantation or Injectable Dosage Form New Animal Drugs; Selenium, Vitamin E Injection

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a previously approved supplemental new animal drug application (NADA) held by Schering-Plough Animal Health Corp. and to remove certain information no longer required in the regulations. The approval concerns use of selenium, vitamin E injection.

EFFECTIVE DATE: May 24, 1999.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209.

SUPPLEMENTARY INFORMATION: Schering-Plough Animal Health Corp., 1095 Morris Ave., P.O. Box 1982, Union, NJ 07083–1982, provided information to support prior approval of supplemental NADA 30–315 for selenium, vitamin E injection. The supplement for use of 2 percent benzyl alcohol instead of 1:10,000 thimerosal had been approved by letter of August 10, 1981. FDA reviewed the information and concurred that the change in ingredient was approved. FDA also reviewed the information requirements of the animal drug regulations and determined that specification of ingredients other than active ingredients is not needed. Therefore, 21 CFR 522.2100 is amended to remove statement of ingredients other than active ingredients.

List of Subjects in 21 CFR Part 522

Animal drugs.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 522 is amended as follows:

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 522 continues to read as follows:

Authority: 21 U.S.C. 360b.

§ 522.2100 [Amended]

2. Section 522.2100 *Selenium, vitamin E injection* is amended in paragraph (a)(1) by removing '', 250 milligrams polyoxyethylated vegetable oil, and 2.0 percent benzyl alcohol, and water for injection''; in paragraph (b)(1) by removing '', 100 milligrams of polyoxyethylated vegetable oil, 1:10,000 thimerosal, and water for injection''; and in paragraphs (c)(1), (d)(1), and (e)(1) by removing '', 250 milligrams polysorbate 80, 2 percent benzyl alcohol, water for injection q.s''.

Dated: May 11, 1999.

Margaret Ann Miller,

Acting Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.

[FR Doc. 99–12963 Filed 5–21–99; 8:45 am]

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DEPARTMENT OF TRANSPORTATION

Coast Guard

33 CFR Part 165

[CGD01–99–038]

RIN 2115–AA97

Safety Zone: Unity Electric Co. Fireworks Display, Shinnecock Bay, Hampton Bays, NY

AGENCY: Coast Guard, DOT.